



Medical Cannabis Cultivation Program

Regulation Development & Environmental Review

Public Scoping, September 2016

The Purpose of Today's Workshop is to...

Provide Information & Obtain Public Input...

Regarding California's medical cannabis cultivation regulations and environmental impacts







Today's Meeting:

- Is Open House Format
- Includes Informational Stations
- Has Technical Experts Available to Discuss Details
- Is an Opportunity to Submit Written or Verbal Comments

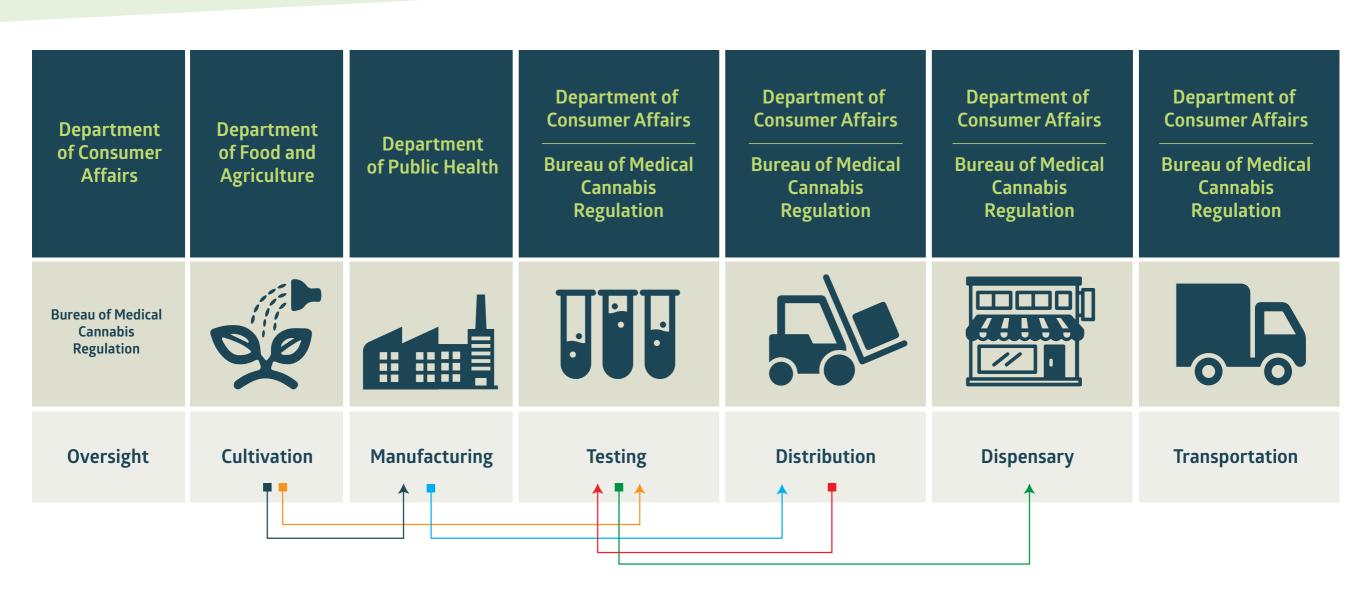


Background

The Medical Cannabis Regulation and Safety Act (MCRSA):

- Signed into law in 2015
- Defined new requirements for medical cannabis production in California
- Established the Medical Cannabis Cultivation Program within the California Department of Food & Agriculture

State Agencies Responsible for Regulating Medical Cannabis Include:



CDFA's Medical Cannabis Cultivation Program (MCCP) will:



License medical cannabis cultivators in the state



Establish conditions under which indoor, outdoor, and mixed-light cultivation may occur



Establish a track and trace program for reporting the movement of medical cannabis items through the distribution chain



Coordinate with other state agencies to protect the environment and human health and safety

CDFA is developing regulations to define statewide licensing requirements for medical cannabis

Licensing will begin on

Jan. 1, 2018,

after environmental review and approval of the regulations by the Office of Administrative Law.



CDFA Will:



Analyze the potential environmental and human health impacts of licensed medical cannabis cultivation



Develop a regulatory structure for statewide medical cannabis cultivation licensing in California



Establish a track and trace system that will document the movement of medical cannabis products from cultivation to final sale

CDFA's medical cannabis cultivation regulations will:

- Establish minimum cultivation licensing requirements for indoor, outdoor, and mixed-light medical cannabis cultivation operations
- Establish a limit on the quantity of licenses issued for cultivation categories over 10,000 square feet
- Ensure that individual and cumulative effects of water diversion and discharge associated with cultivation do not affect instream flows needed for fish and natural flows
- Ensure that cultivation will not negatively impact springs, riparian wetlands, and aquatic habitats

CDFA's medical cannabis cultivation regulations will:

- Require that licensed cannabis cultivation is conducted in accordance with state and local laws
- Establish procedures for the issuance, reporting and revocation of medical cannabis cultivation licenses
- Establish a scale of application, licensing, and renewal fees, based upon the cost of administering and enforcing the Program
- Develop a cultivation checklist tool to evaluate environmental impacts of cannabis cultivation licensing

CDFA is authorized to issue ten license types for medical cannabis cultivation

License Type	
Type 1	Specialty Outdoor
Type 1A	Specialty Indoor
Type 1B	Specialty Mixed-Light
Type 2	Small Outdoor
Type 2A	Small Indoor
Type 2B	Small Mixed-Light
Type 3	Outdoor
Type 3A	Indoor
Type 3B	Mixed-Light
Type 4	Nursery

CDFA will not begin issuing licenses until 2018

CDFA's Regulations will Include Guidelines on:

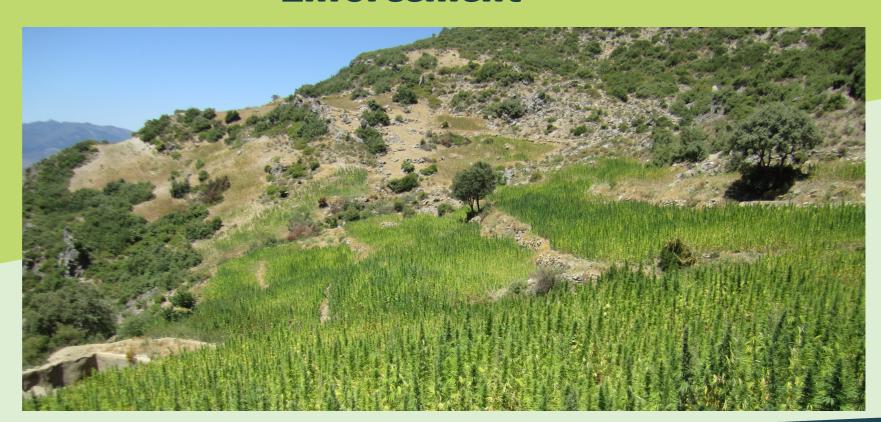
- Applying for a Cultivation
 License
- Cultivator Requirements
- Track and Trace Requirements
- Inspections
- Enforcement



Key Considerations for Developing Medical Cannabis Cultivation Regulations Include:

- Pesticide, Fertilizer, and other Chemical Use
- Energy Use
- Water Use
- Cultivation

- Transportation & Tracking
- Other MCRSA licensing programs
- Enforcement



CDFA is preparing a Programmatic Environmental Impact Report (PEIR) for Statewide Medical Cannabis Cultivation to:

- Ensure that medical cannabis cultivation is protective of the environment, cannabis cultivation workers, and the general public
- Establish a track and trace program to ensure the movement of medical cannabis is tracked throughout the production chain

The primary purpose of CDFA's Programmatic Environmental Impact Report (PEIR) is to:



Analyze and disclose the **reasonably foreseeable environmental impacts** that may occur as a result of the Medical Cannabis Cultivation Program (MCCP)



Analyze and disclose the **potentially significant environmental impacts** associated with the MCCP



Where impacts are significant, identify potentially feasible mitigation measures and alternatives that substantially lessen or avoid such effects



Potential environmental issues to be addressed in the Programmatic Environmental Impact Report (PEIR) include:

- Aesthetics
- Agriculture and Forestry Resources
- Air Quality
- Biological Resources
- Cultural Resources
- Geology and Soils
- Greenhouse Gas Emissions
- Hazards and Hazardous Materials
- Hydrology and Water Quality

- Land Use and Planning
- Mineral Resources
- Noise
- Population and Housing
- Public Services
- Recreation
- Transportation and Traffic
- Tribal Cultural Resources
- Utilities and Service Systems
- Cumulative Impacts
- Irreversible Impacts

CDFA Wants Your Input On...



Potential Environmental Impacts of Medical Cannabis Cultivation



Scope of the Proposed Regulations



Reasonable Range of Alternatives



Methodologies for Impact Analysis



Types of Impacts to Evaluate



Potential Mitigation Strategies

The Public Comment Period is from September 1 – 30, 2016

Next Steps





Public Scoping Comment Period

September 2016



MCCP Regulation Development

Fall 2016



Environmental Review



Proposed Regulation to Office of Administrative Law (OAL)

Early 2017



Draft Programmatic Environmental Impact Report

Spring 2017



Public Review and Comment

Spring 2017



Public Review and Comment

Spring 2017



Final Regulation to Office of Administrative Law (OAL)

Fall 2017



Final PEIR & Notice of Determination

Winter 2017



License Issuance

January 2018





Comment Period: September 1-30, 2016



California Department of Food and Agriculture Attn: Amber Morris Medical Cannabis Cultivation Comments 1220 N Street, Suite 400 Sacramento, CA 95814

@ EMAIL TO:

mccp.peir@cdfa.ca.gov

AT PUBLIC MEETING:



Fill out a comment form or give verbal comments to a court reporter.





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This Concludes the Presentation

Please speak to staff for additional information and questions

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